Lys-D-Pro-Thr

| Cat. No.: | HY-125083 | |
|----------------------|---|---|
| CAS No.: | 117027-34-6 | ОН ОН |
| Molecular Formula: | $C_{15}H_{28}N_4O_5$ | 0 |
| Molecular Weight: | 344.41 | |
| Sequence: | Lys-{d-Pro}-Thr | $H_{2}N_{2} \wedge A_{2} = \int_{-\infty}^{0} \int_{-\infty}^{0}$ |
| Sequence Shortening: | L-{d-Pro}-T | |
| Target: | Interleukin Related | |
| Pathway: | Immunology/Inflammation | |
| Storage: | Sealed storage, away from moisture | |
| | Powder -80°C 2 years | |
| | -20°C 1 year | |
| | * In solvent : -80°C, 6 months; -20°C, 1 month (sealed storage, away from moisture) | |

SOLVENT & SOLUBILITY

| In Vitro | DMSO : 100 mg/mL (290.35 mM; Need ultrasonic) | | | | | |
|----------|---|---|---|------------|------------|--|
| | | Solvent Mass Concentration | 1 mg | 5 mg | 10 mg | |
| Pre | Preparing Stock Solutions | 1 mM | 2.9035 mL | 14.5176 mL | 29.0352 mL | |
| | 5 mM 0.5807 mL 2.9035 m 10 mM 0.2904 mL 1.4518 m | 2.9035 mL | 5.8070 mL | | | |
| | | 10 mM | 0.2904 mL | 1.4518 mL | 2.9035 mL | |
| | Please refer to the solubility information to select the appropriate solvent. | | | | | |
| In Vivo | 1. Add each solvent o Solubility: ≥ 2.5 mg | ne by one: 10% DMSO >> 40% PE ;/mL (7.26 mM); Clear solution | 2: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline 5 mM); Clear solution | | | |
| | 2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.5 mg/mL (7.26 mM); Clear solution | | | | | |
| | 3. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.5 mg/mL (7.26 mM); Clear solution | | | | | |

| BIOLOGICAL ACTIV | |
|---------------------------|--|
| | |
| Description | Lys-D-Pro-Thr, an IL-1beta analogue, is a potent IL-1 inhibitor. Lys-D-Pro-Thr inhibits the protec |
| IC ₅₀ & Target | IL-1 |
| | |
| In Vivo | Lys-D-Pro-Thr (10 mg/kg; IP; for 4 days) inhibits the protective effect of fMLP (HY-P0224) and the |



| MCE has not independe | ently confirmed the accuracy of these methods. They are for reference only. |
|-----------------------|---|
| Animal Model: | 11-d-old rats ^[1] |
| Dosage: | 10 mg/kg |
| Administration: | IP; for 4 days |
| Result: | Inhibited the protective effect of fMLP and that of MMK-1 against Etoposide-induced alopecia. It alone did not induce alopecia. |

REFERENCES

[1]. Takahiro Tsuruki, et al. Mechanism of the protective effect of intraperitoneally administered agonists for formyl peptide receptors against chemotherapy-induced alopecia. Biosci Biotechnol Biochem. 2007 May;71(5):1198-202.

[2]. Y Uehara, et al. Central administration of Lys-D-Pro-Thr, an interleukin-1 beta 193-195 analogue, stimulates feeding in rats. Neuropeptides. 1991 May;19(1):9-11.

Caution: Product has not been fully validated for medical applications. For research use only.